



State of Vermont
Department of Vermont Health Access
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Agency of Human Services

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REQUEST FOR AIRWAY CLEARANCE DEVICES

PLEASE COMPLETE BOTH EVALUATION SEGMENTS DURING THE TRIAL

PLEASE SUBMIT COMPLETED FORM TO DME PROVIDER

Patient Name: _____ Date of Birth: ____/____/_____
Member ID#: _____ Home Phone #: _____ Cell Phone #: _____ Work Phone #: _____
Patient Address: _____ City, State, Zip code: _____

Physician Name: _____ Physician NPI: _____ Medicaid Provider Number: _____
Physician Address: _____ City, State, Zip code: _____
Physician Phone #: _____ Physician Fax#: _____
Physician Signature: _____

Durable Medical Equipment (DME) Requested:

DME Description: _____ HCPCS code: _____
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<u>For Airway Oscillating Devices and Mechanical Percussors:</u> 1.Does the patient have cystic fibrosis/CF, chronic bronchitis, bronchiectasis, immotile cilia syndrome, or asthma?	<input type="checkbox"/> Yes <input type="checkbox"/> No Comments: _____
<u>For Positive Expiratory Pressure/PEP Masks:</u> 1.Does the patient have CF, chronic bronchitis, immotile cilia syndrome, asthma, or chronic obstructive pulmonary disease/COPD?	<input type="checkbox"/> Yes <input type="checkbox"/> No Comments: _____
<u>For High-Frequency Chest Compression Systems:</u> 1.Has the patient failed standard treatments to adequately mobilize retained secretions? 2.Does the patient have bronchiectasis confirmed by CT scan characterized by daily productive cough for at least 6 months or by frequent (more than 2 times per year) exacerbations requiring antibiotic therapy? 3.Does the patient have cystic fibrosis or immotile cilia syndrome? 4.Is the patient within the first 6 months post-operatively following lung transplant and unable to tolerate standard chest physiotherapy?	<input type="checkbox"/> Yes <input type="checkbox"/> No Comments: _____ <input type="checkbox"/> Yes <input type="checkbox"/> No Comments: _____ <input type="checkbox"/> Yes <input type="checkbox"/> No Comments: _____ <input type="checkbox"/> Yes <input type="checkbox"/> No Comments: _____

5.Does the patient have one of the following neuromuscular diseases?	<input type="checkbox"/> Acid maltase deficiency <input type="checkbox"/> Hereditary muscular dystrophy <input type="checkbox"/> Anterior horn cell disease, including amyotrophic lateral sclerosis <input type="checkbox"/> Multiple Sclerosis <input type="checkbox"/> Myotonic disorder <input type="checkbox"/> Paralysis of the diaphragm <input type="checkbox"/> Post-polio <input type="checkbox"/> Quadriplegia <input type="checkbox"/> Other myopathies
<u>For Mechanical In-Exsufflation Devices:</u> 1.Does the patient have a neuromuscular disease that is causing a significant impairment of chest wall and/or diaphragmatic movement and for whom standard treatments have not been successful in adequately mobilizing retained secretions?	<input type="checkbox"/> Yes <input type="checkbox"/> No Comments:
<u>There must be well-documented failure of standard treatments to adequately mobilize retained secretions:</u> Document trial/consideration of each applicable device/technique trialed listed below, AND why it was not successful for the beneficiary. <input type="checkbox"/> CPT (Manual or Percussor) <input type="checkbox"/> PEP <input type="checkbox"/> Flutter/Acapella <input type="checkbox"/> Cough Assist <input type="checkbox"/> Breathing/Drainage Techniques <input type="checkbox"/> No Caregiver Available <input type="checkbox"/> Physical Limitations of Caregiver <input type="checkbox"/> GERD <input type="checkbox"/> Physical Limitations of Patient <input type="checkbox"/> Did not Mobilize Secretions <input type="checkbox"/> Young Age <input type="checkbox"/> Too Fragile for Percussion <input type="checkbox"/> Resistance to Therapy <input type="checkbox"/> Aspiration Risk <input type="checkbox"/> Can't Tolerate Positioning <input type="checkbox"/> Insufficient Expiratory Force <input type="checkbox"/> Artificial Airway <input type="checkbox"/> Severe Arthritis/Osteoporosis <input type="checkbox"/> Kyphosis/Scoliosis <input type="checkbox"/> Cognitive Level <input type="checkbox"/> Spasticity/Contractures <input type="checkbox"/> Inability to Form Mouth Seal <input type="checkbox"/> Other _____ Reason for failure:	

Trial Period- item requested must have a 3-month trial period.	First Evaluation: Baseline ____/____/____	Final Evaluation (at least 3 months from first): final determination of device efficacy ____/____/____
Prescription for device use (include minutes/day):		
Hospitalizations including dates and reason for admission or IV antibiotic therapy including dates and reason:		
Requires assistance to mobilize:	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
Adequate physiological cough reflex:	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
Pulmonary Functions: (please attach results)	Pre use of device:	With device use:
Daily productive cough: Duration in months:	<input type="checkbox"/> Yes <input type="checkbox"/> No _____	<input type="checkbox"/> Yes <input type="checkbox"/> No _____
Is member ventilator dependent:	<input type="checkbox"/> Yes <input type="checkbox"/> No	<input type="checkbox"/> Yes <input type="checkbox"/> No
Is there assistance of a caregiver in the home:	<input type="checkbox"/> Yes <input type="checkbox"/> No	<input type="checkbox"/> Yes <input type="checkbox"/> No
Documented adherence to therapy is required: (please attach data chart)	<input type="checkbox"/> Yes <input type="checkbox"/> No	<input type="checkbox"/> Yes <input type="checkbox"/> No
Education provided to member and family if applicable:	<input type="checkbox"/> Yes <input type="checkbox"/> No	<input type="checkbox"/> Yes <input type="checkbox"/> No
Specify brands trialed:		
Other Pertinent Information:		
MD Signature:		